4339 FIRST AID KIT- 4339 first aid kit Honeywell Safety Products USA, INC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

4339 First Aid Kit (1st aid Sp, EW, aypanal EX, miralac, amm. Inh, triple, burn spray WS, alcohol wipe, BZK wipes - Z019829)

Eyewas h Active ingredient

Sterile Water 99%

Eyewash *Purpose*

Eyewash

Eyewash

Uses

• for flushing the eye to remove loose foreign material, air pollutants or chlorinated water

Eyewash

Warnings

For external use only Obtain immediate medical treatment for all open wounds in or near eyes. To avoid contamination, do not touch tip of container to any surface. Do not reuse. Once opened, discard.

Do not use

- if solution changes color or becomes cloudy
- if you have open wounds in or near the eyes, get medical help right away.

Stop use and ask a doctor if

- you experience eye pain
- changes in vision
- continued redness or irritation of the eye
- condition worsens or persists

Keep out of reach of children

• If swallowed, get medical help or contact a Poison Control Center right away.

Evewash

Directions

- remove contacts before using
- twist top to remove
- flush the affected area as needed
- control rate of flow by pressure on the bottle
- if necessary, continue flushing with emergency eyewash or shower

Eyewash

Inactive ingredients

sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic

Eyewash Questions

1-800-430-5490

Ammonia Active ingredient

Ammonia 15%

Ammonia

Purpose

Respiratory stimulant

Ammonia

Uses

• to prevent or treat fainting

Ammonia

Warnings

For external use only

Do not use

• if you have breathing problems such as asthma or emphysema

Stop use and ask a doctor if

• condition persists

Keep out of reach of children

• If swallowed get medical help or contact a Poison Control Center right away.

Ammonia

Directions

- hold inhalant away from face and crush ampoule between thumb and forefinger at position indicated on sleeve.
- hold near nostrils for inhalation of volatile vapor

Ammonia

Other information

• store at room temperature away from light

Ammonia

Inactive ingredient

alcohol USP, FD&C red #40, lavender oil, lemon oil fcc, nutmeg oil, purified water

Ammonia

Questions or Comments?

1-800-430-5490

Water Soluble 1st Aid Spray Active ingredient

Benzethonium chloride 0.2% w/w - Benzocaine 10% w/w

Water Soluble 1st Aid Spray *Purpose*

Topical antiseptic

Topical anesthetic

Water Soluble 1st Aid Spray

Uses

for temporary relief of pain and itching and helps protect against infection in

- minor cuts and scrapes
- insect bites
- minor skin irritations

Water Soluble 1st Aid Spray Warnings

For external use only

Flammable

- keep away from fire or flame
- contents under pressure
- do not puncture or incinerate container
- do not expose to temperature above 120 0 F

Do not use

- in the eyes or other mucous membranes
- in cases of serious burns
- in case of deep orpuncture wounds
- for a prolonged period of time
- on large portion of the body

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days
- condition clears up and occurs again within a few days
- redness, swelling, or irritation occurs

Keep out of reach of children

• If swallowed, get medical help or contact a Poison Control Center right away

Water Soluble 1st Aid Spray Directions

- clean the affected area
- shake can well before using
- hold 4 6 inches from surface and spray area until wet
- may be covered with a sterile bandage. If bandaged, let dry first
- for adult institutional use only
- not intended for use on children

Water Soluble 1st Aid pray Other information

- avoid inhaling
- use only as directed
- intentional misuse by deliberately concentrating and inhaling the contents may be harmful or fatal

Water Soluble 1st Aid pray Inactive ingredients

dipropylene glycol, isobutane, N-butane, propane

Burn Relief Water Soluble Active ingredients

Benzethonium chloride 0.2% w/w

Benzocaine 10% w/w

Menthol 0.33% w/w

Burn Relief Water Soluble Purpose

Topical antiseptic

Topical anesthetic

Topical anesthetic

Burn Relief Water Soluble

Uses

for the temporary relief of pain and itching and helps protect against infection in:

- minor cuts and scrapes
- burns
- sunburn
- insect bites
- minor skin irritations

Burn Relief Water Soluble Warnings

For external use only

Flammable keep away from fire or flame

- contents under pressure
- do not puncture or incinerate container
- do not expose to temperatures above 120 0 F

Do not use

- in or near the eyes or other mucous membranes
- in case of serious burns
- in case of deep or puncture wounds
- for prolonged period of time
- on large portion of the body

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days
- condition clears up and recurs within a few days
- redness, swelling, or irritation occurs

Keep out of the reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Burn Relief Water Soluble

Directions

- clean the affected area
- shake can well before using
- hold 4 6 inches from surface and spray area until wet
- may be covered with a sterile bandage, if bandaged let dry first
- for adult institutional use only
- not intended for use on children

Burn Relief Water Soluble

Other information

- avoid inhaling
- use only as directed
- intentional misuse by deliberately concentrating or inhaling the contents may be harmful or fatal

Burn Relief Water Soluble

Inactive ingredients

dipropylene glycol, isobutane, n-butane, propane

Triple

Active ingredients

Bacitracin zinc 400 units

Neomycin sulfate (5 mg equivalent to 3.5 mg Neomycin base)

Polymyxin B sulfate 5000 units

Triple *Purpose*

First aid antibiotic

First aid antibiotic

First aid antibiotic

Triple

Uses

- first aid to help prevent infection in
- minor cuts
- scrapes
- burns

Triple

Warnings

For external use only

Allergy alert do not use if you are allergic to any of the ingredients

Do not use

- in the eyes
- over large areas of the body

Ask a doctor before use if you have

- a deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- the condition persists or gets worse
- a rash or other allergic reaction develops
- you need to use longer than 1 week

Keep out of the reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Triple

Directions

- clean the affected area
- apply a small amount of the product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily
- may be covered with a sterile bandage

Triple

Other information

- store at 15 0 to 25 0 C (59 0 to 77 0 F)
- tamper evident sealed packets

• do not use if packet is torn or opened

Triple

Inactive ingredient

petrolatum

Alcohol

Active ingredient

Isopropyl alcohol 70%

Alcohol

Purpose

First aid antiseptic

Alcohol

Uses

• first aid to help prevent infection in minor cuts, scrapes, and burns

Alcohol

Warnings

For external use only

Flammable, keep away from fire or flame.

Do not use

- in the eyes
- over large areas of the body

Ask a doctor before use if you have

deep or puncture wounds

animal bites

serious burns

When using this product

• do not use longer than one week unless directed by a doctor

Stop use and consult a doctor

• if condition persists or gets worse

Keep out of reach of children

• If swallowed, get medical help or contact a Poison Control Center right away.

Alcohol

Directions

- clean the affected area
- apply wipe to affected area 1 to 3 times daily

- may be covered with a sterile bandage
- discard wipe after single use

Alcohol

Other information

store at room temperature 15 0 to 25 0 C (59 0 to 77 0 F)

Alcohol

Inactive ingredient

water

Miralac

Active ingredient (in each chewable tablet)

Miralac

Active ingredient (in each chewable tablet)

Calcium carbonate 420 mg

Miralac

Purpose

Antacid

Miralac

Uses

for the relief of

- acid indigestion
- heartburn
- sour stomach
- upset stomach associated with these symptoms

Warnings

Ask a doctor before use if you have

- kidney stones
- calcium-restricted diet

Ask a doctor before use if you are

• presently taking a prescription drug. Antacids may interfere with certain prescription drugs

When using this product

• do not take more than 12 tablets in a 24- hour period, or use the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a doctor.

Keep out of the reach of children.

In case of accidental overdose, seek professional assistance or contact a Poison Control Center immediately.

Miralac

Directions

• chew 1 to 2 tablets every 4 hours as symptoms occur, or as directed by a doctor.

Miralac

Other information

- each tablet contains: calcium 170 mg
- sucrose free
- lactose free
- store at room temperature
- TAMPER EVIDENT PACKETS- DO NOT USE IF OPEN OR TORN

Miralac

Inactive ingredients

magnesium stearate, mint flavor, silicon dioxide, sorbitol, starch

Miralac

Questions or comments?

1-800-430-5490

BZK

Active ingredient

Benzalkonium chloride 0.13% w/v

BZK

Purpose

First aid antiseptic

BZK

Uses

Antiseptic cleansing of face, hands, and body without soap and water

BZK

Warnings

For external use only

Do not use

- in the eyes or over large areas of the body
- on mucous membranes
- on irritated skin
- in case of deep puncture wounds, animal bites or serious burns, consult a doctor
- longer than 1 week unless directed by a doctor

Stop use and ask a doctor if

- if irritation, redness or other symptoms develop
- the condition persists or gets worse

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

BZK

Directions

tear open packet and use as a washcloth

BZK

Other information

- store at room temperature 15 0 to 30 0 C (5 0 86 0 F)
- do not reuse towelette

BZK

Inactiave ingredient

water

BzK

Questions

1-800-430-5490

Aypanal

Active ingredient (in each tablet)

Acetaminophen 500 mg

Aypanal

Purpose

Pain reliever/fever reducer

Aypanal

Uses

- temporarily relieves minor aches and pains due to the common cold and headache
- temporarily reduces fever

Aypanal

Warnings

Liver Warning: This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 4,000 mg in 24 hours, which is the maximum daily amount.
- with other drugs containing acetaminophen
- adult has 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If skin reaction occurs, stop use and seek medical help right away

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription).
- If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

Ask a doctor before use if you have

• liver disease

Ask a doctor or pharmacist before use if

• you are taking the blood thinning drug warfarin

Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- new symptoms occur
- redness or swelling is present

If pregnant or breastfeeding

• ask a health professional before use.

Keep out of reach of children.

Keep out of reach of children.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Aypanal

Directions

- do not take more than directed (see overdose warning)
- adults and children 12 years of age and over: Take 2 tablets with water every 6 hours while symptoms last.
- do not take any more than 8 tablets in 24 hours.
- children under 12: consult a doctor

Aypanal

Other information

- store at room temperature 15 0 -30 0 C (59 0 -86 0 F)
- TAMPER EVIDENT- DO NOT USE IF OPEN OR TORN

Aypanal

Inactive igredients

microcrystalline cellulose, povidone, sodium starch glycolate, starch, stearic acid

Aypanal Questions or Comments

1-800-430-5490

4339 Z019829 KIT CONTENTS

- 1 1X3 PLASTIC 100/BOX
- 1 FINGERTIP "T" WOVEN 40/BOX
- 1 1X3 WOVEN SING 50/BOX
- 1 SWIFT KNUCKLE 40/BX
- 1 AMMONIA INHALANTS 10 PER
- 1 EYE DRESS PKT W/4 ADH STRIPS
- 1 TWEEZER PLASTICS 4"
- 1 O/H TAPE ADHESIVE TRI-CUT
- 1 FIRST AID GUIDE ASHI
- 1 BLOODSTOPPER
- 10 NON-ADHERENT PADS 2"X3" 10'S
- 1 GZE PADS STERILE 3"X 3" 25'S
- 3 CTA 3" SINGLE TIP 100/PER
- 1 ANTISEPTIC WIPES BZK CHL 20'S
- 1 FIRST AID SPRAY AEROSOL 3 OZ
- 1 ALCOHOL WIPES 50'S
- 1 AYPANAL EX-STR 2/ENV 250
- 1 MIRALAC TABS IND PK 2/ENV 250
- 1 BURN SPRAY 3 OZ
- 1 TRIPLE BIOTIC .5 GRAM PKT 20
- 1 4OZ BFS EYEWASH TRILINGUAL BOTTLE
- 1 SCISSOR BDGE 4" RED PLS HDL
- 1 LBL STOCK 6-3/8"X4"
- 1 LBL STOCK 4"X2-7/8"
- 1 LBL CONTS 8"X8", CUSTOM ID B
- 1 LABEL COVER, GRAINGER Z019829
- 2 PR LRG NITRILE GLVES
- 1 KIT ST MED FA CABINET-SP SHELF
- 1 TRI BNDG NON WOVEN 40"X40"X56"
- 1 COLD PACK UNIT 4"X6" BULK

Eyewash

Solución

Isotónico Estéril

Isotonic Solution Isotonique Stérile 16 fl. oz. (473 mL)

La Solution

Sterile

Drug Facts (for USA only) Active ingredient Uses
for flushing the eye to remove loose foreign material, air pollutants, or chlorinated water. or chlorinated water. Warnings
For external use only - Obtain immediate medical treatment for all open wounds in or near the eyes. To avoid contamination, do not buck by of container to any surface. Do not reuse. Once opened, discard. Do not use
• if solution changes color or becomes cloudy
• if you have open wounds in or near the eyes, get medical help Stop use and consult a doctor if: you experience eye pain
 ontinued redness or irritation of the eye
 condition worsens or persists condition worsens or persists
 Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Directions

• remove contacts before using
• twist top to remove
• flush the affected area as needed
• control rate of flow by pressure on the bottle
• if necessary, continue flushing with emergency eyewash or shower Inactive ingredients sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic Questions? Call 1-800-430-5490 Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

and supplier

#32-000454

NUEVO PEDIDO.

REORDER

#32-004510 Rev. J

LABEL

Purpose Eyewash

Datos de medicamento (Para EE.UU. solamente) Usos para el lavado de ojo para quitar las particulas sueltas y extrañas, los contaminantes aeros, o agua de cloruro Advertencias

Para el uso externo sólo - Obtenga tratamiento médico inmediato para todas las heridas abiertas en o cerca de los ojos. Para evitar la contaminación, no toque la punta del envase con inguna superficie. No vuedva a usar. Vez abierto, descarte. No se use • si la solución se enturbia o cambia de color • si tiene heridas abiertas en o cerca del ojo, obtenga ayuda medica de inmediato de inmediato
Deje de usar y consulte a un médico si:

experimenta dolor de ojo

cambio de visión

rojez continuo o irritación del ojo

la condición empeora o persiste Manténgase fuera del alcance de los niños.
En caso de ingestión accidental, obtenga atención médica o llame a un Centro de control de envenenamiento inmediatamente. Instrucciones

quitese los lentes de contacto antes de usar la solución

tuerza la tapa para quitar

en juague el área afectada según se necesite

controle el chorro haciendo presión el la botella

si es necesario, sique enjuagado con un lavaojos o ducha de emergencia Ingredientes inactivos cloruro de sodio, fosfato de sodio dibásico, fosfato de sodio monobási ¿Preguntas? Llame al 1-800-430-5490 Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

Information

Usages
Pour le rinçage des yeux afin d'enlever un corps étranger, des polluants atmospheriques ou de l'eau chlorée

Advertissements
Pour usage externe seulement - Obtenir immédiatement des
soins médicaux pour toutes les plaies ouvertes dans ou près des
yeux. Pour éviter toute contamination, ne pas toucher la pointe du
récipient à n'importe quelle surface. Ne pas réutiliser. Une fois
ouvert, jetez-les.
Ne pas utilité.

ouvert, jetez-les.

Ne pas utiliser

• si la solution a changé de couleur ou si elle est brouillée

• si la solution a changé de couleur ou si elle est brouillée

• si vous avez des plaies ouvertes aux yeux ou à proximité,
consultez immédiatement un médecin

Cesser d'utiliser la solution et consulter un médecin

• vous ressentez une douleur oculaire

• si votre vision change

• rougeur ou imfation persistante des yeux

• condition empire ou persiste

Garder hors de la portée des enfants.

En cas d'ingestion, communiquer immédiatement avec un médecin

ou avec un centre antipoison.

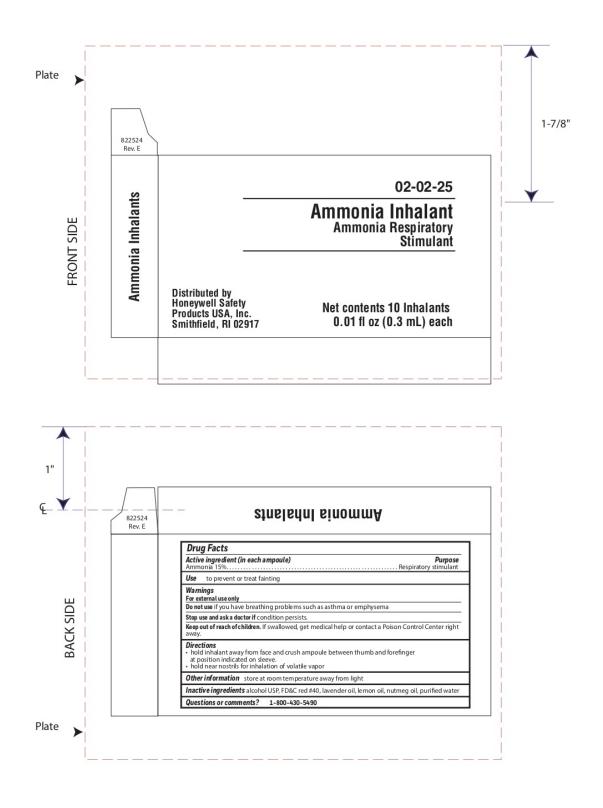
Mode d'emploi

• enlever les verres de contact avant l'utilisation • dévisser le bouchon pour l'enlever • rincer la zone touchée selon les besoins • ajuster le débit d'écoulement de la solution en se solution en contenar et inécessaire, confluirer de rincer avec unesolution de rinçage oculaire d'urgence ou une douche

Ingrédients eau stérile, chlorure de sodium, phosphate dibasique de sodium, phosphate monobasique de sodium

Des questions? Faites le 1-800-430-5490 Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

Ammonia Principal Display Panel



796006 Rev. E (page 3 of 3)

SHAKE WELL BEFORE USING

Honeywell **FIRST AID** ANTISEPTIC SPRAY

Water soluble Benzethonium chloride Topical antiseptic

Benzocaine

Topical anesthetic

Helps prevent infection and relieves pain.

CAUTION: FLAMMABLE

Contents under pressure Read warning on back panel.

NET WT. 3 OZ (85gm.)

Cat. No. 151019

DRUG FACTS

Active ingredients

Purpose

Benzethonium chloride 0.2% w/w. Benzocaine 10% w/w.

.Topicalantiseptic

• for the temporary relief of pain and itching and helps to protect against infection in

 minor cuts and scrapes · insect bites minor skin irritations

Warnings For external use only

Flammable • keep away from fire or flame

 do not puncture or incinerate container do not expose to temperatures above 120°F

Do not use • in or near eyes or other mucus membranes • in case of serious burns
• in case of deep or puncture wounds • for a prolonged period of time

on large portion of the body

Stop use and ask a doctor if:

conditions worsens or symptoms persist for more than 7 days

condition clears up and recurs within a few days
 redness, swelling or irritation occurs

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions • clean the affected area • shake can well before using

hold 4-6 inches from surface and spray area until wet
 may be covered with a sterile bandage. If bandaged, let dry first
 for adult institutional use only
 not intended for use on children

Other information avoid inhaling · use only as directed

intentional misuse by deliberately concentrating and inhaling the contents may be harmful

Inactive ingredients dipropylene glycol, isobutane, n-butane, propane

Questions or comments? 1-800-430-5490



Distributed by Honeywell Safety Products USA, Inc Smithfield, RI 02917

Honeywell

Burn Relief Water Soluble Principal Display Panel

SHAKE WELL BEFORE USING

Honeywell BURN SPRAY

Water soluble **Benzethonium chloride**

Topical antiseptic

Benzocaine Topical anesthetic

Menthol

Topical anesthetic

Provides antiseptic treatment and helps relieve the pain of minor burns and sunburn.

CAUTION: FLAMMABLE

Contents under pressure Read warning on back panel.

NET WT. 3 OZ (85gm.)

Cat. No. 201005

DRUG FACTS

Active ingredients

Benzethonium chloride 0.2% w/w. Benzocaine 10% w/w... Menthol .33%

Purpose Topical antiseptic Topical anesthetic Topical anesthetic

Uses • for the temporary relief of pain and itching and helps to protect against infection in · minor cuts and scrapes · burns · sunburn · insect bites · minor skin i rritations

Warnings

For external use only

Flammable • keep away from fire or flame • contents under pressure

do not puncture or incinerate container do not expose to temperatures above 120°F Do not use • in or near eyes or other mucus membranes • in case of serious burns

in case of deep or puncture wounds
 for a prolonged period of time

· on large portion of the body

Stop use and ask a doctor if:

- · conditions worsens or symptoms persist for more than 7 days
- · condition clears up and recurs within a few days

redness, swelling or irritation occurs

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions • clean the affected area • shake can well before using

- hold 4-6 inches from surface and spray area until wet
 may be covered with a sterile bandage. If bandaged, let dry first
- for adult institutional use only
 not intended for use on children

Other information avoid inhaling · use only as directed

intentional misuse by deliberately concentrating and inhaling the contents may be harmful

Inactive ingredients dipropylene glycol, isobutane, n-butane, propane

Questions or comments? 1-800-430-5490

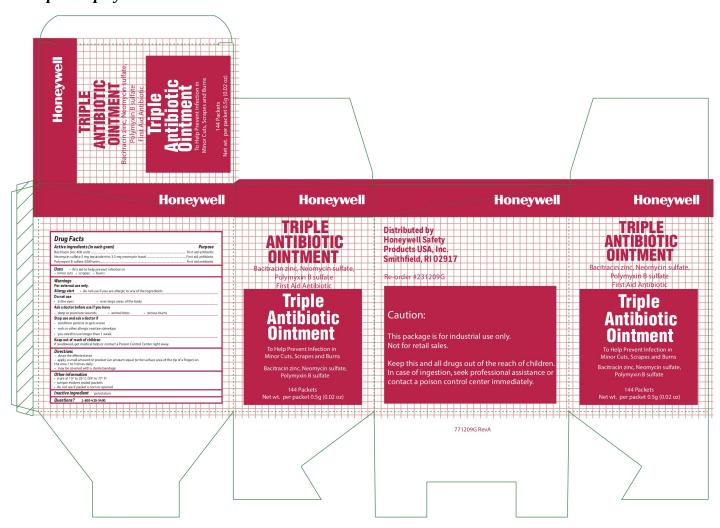


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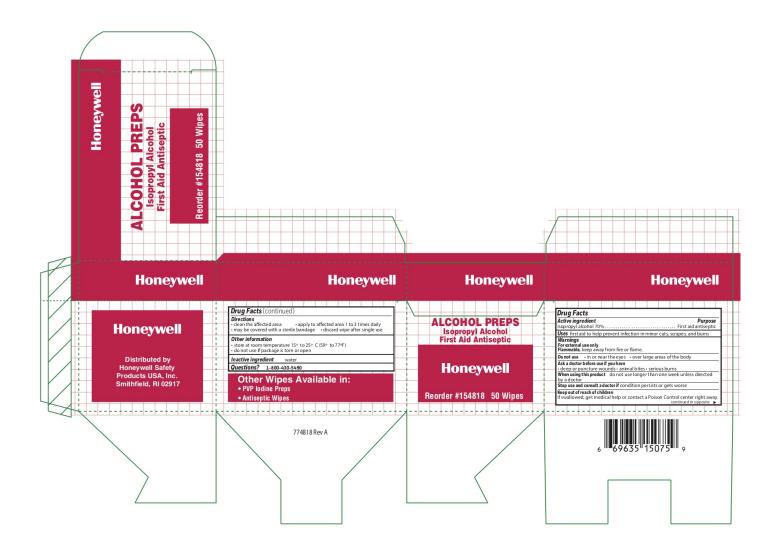
Honeywell

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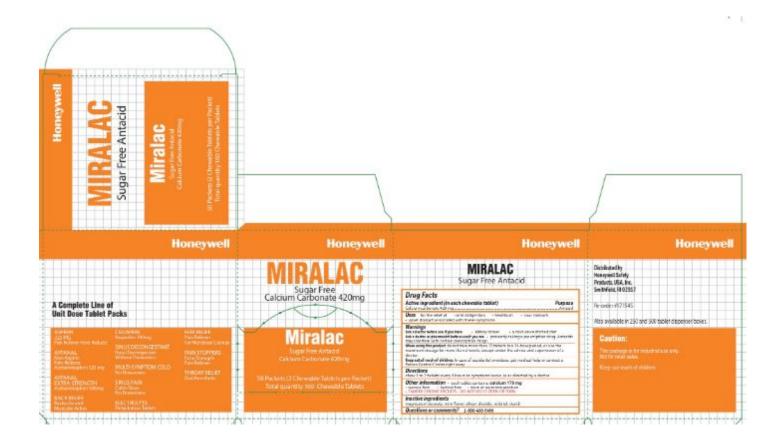
Triple
Principal Display Panel



Alcohol Principal Display Panel

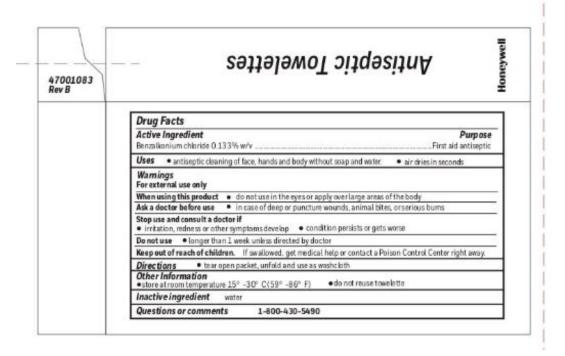


Miralac Principal Display Panel

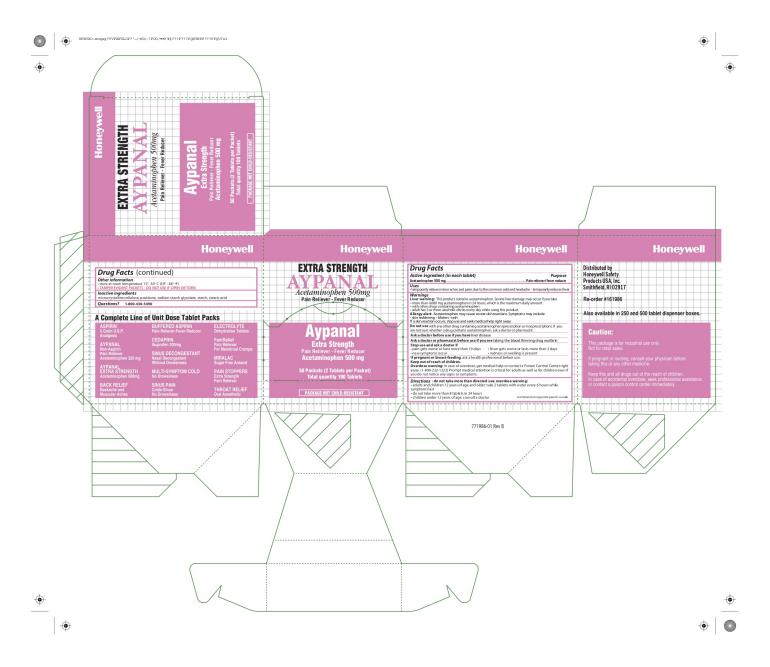


BZK Principal Display Panel

	Honeywell	S
02-16-35MD		lette
Antiseptic Towelette	-	оме
Benzalkonium chlorio First aid antisept		Antiseptic Towelettes
Six-Saturated Towelette	10	tise
	Distributed by Honeywell Safety Products USA, Inc. Smithfield, RI 02917	An



Aypanal Principal Display Panel



4339 Kit Label Z019829

CABINET, BULK 150 PERSON







GRAINGER.COM®

Distributed by Honeywell Safety Products USA, Inc. Smithfield, RI 02917

4339 FIRST AID KIT

4339 first aid kit kit

Product Information

HUMAN OTC DRUG NDC:0498-4339 Product Type Item Code (Source)

Packaging

ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:0498-4339-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	118 mL
Part 2	10 AMPULE	3 mL
Part 3	1 CAN	85 g
Part 4	1 CAN	85 g
Part 5	20 PACKET	10 g
Part 6	50 POUCH	20 mL

Part 7	125 PACKET	250
Part 8	20 PACKET	28 mL
Part 9	125 PACKET	250

Part 1 of 9

EYESALINE EMERGENCY EYEWASH

purified water liquid

Product Information

Item Code (Source)NDC:0498-0100Route of AdministrationOPHTHALMIC

Active Ingredient/Active Moiety Ingredient Name Basis of Strength WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R) WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)

Inactive Ingredients	
Ingredient Name	Strength
SO DIUM PHO SPHATE, MO NO BASIC, MO NO HYDRATE (UNII: 593YOG76RN)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SO DIUM PHO SPHATE, DIBASIC (UNII: GR686LBA74)	

l	Packaging				
l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NI	OC:0498-0100-02	118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Info	rmation		
Marketing Category Application Number or Monograph Citation Marketing Start Dat			Marketing End Date
OTC monograph final	part349	12/18/20 18	

Part 2 of 9

AMMONIA INHALENT

ammonia inhalent inhalant

Product Information

Item Code (Source)	NDC:0498-3334
Route of Administration	RESPIRATORY (INHALATION)

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength Strength	
AMMO NIA (UNII: 5138 Q 19 F1X) (AMMO NIA - UNII:5138 Q 19 F1X)	AMMONIA	0.045 g in 0.3 mL

Inactive Ingredients	
Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	

l	Packaging				
l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1	NDC:0498-3334-00	0.3 mL in 1 AMPULE; Type 0: Not a Combination Product		

Marketing Info	rmation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/18/2018	

Part 3 of 9

FIRST AID ANTISEPTIC WATER SOLUBLE

benzethonium chloride, benzocaine spray

Product Information	
Item Code (Source)	NDC:0498-0031
Route of Administration	TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZETHO NIUM CHLO RIDE (UNII: PH41D05744) (BENZETHO NIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 g in 100 g	
BENZO CAINE (UNII: U3RS Y48 JW5) (BENZO CAINE - UNII: U3RS Y48 JW5)	BENZOCAINE	10 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
ISOBUTANE (UNII: BXR49TP611)		
BUTANE (UNII: 6LV4FOR43R)		

PROPANE (UNII: T75W9911L6)

DIPROPYLENE GLYCOL (UNII: E107L85C40)

l	Packaging			
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:0498-0031-40	85 g in 1 CAN; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/19/2018	

Part 4 of 9

BURN WATER SOLUBLE

benzocaine, benzethonium chloride, menthol spray

Product Information	
Item Code (Source)	NDC:0498-0021
Route of Administration	TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZETHO NIUM CHLO RIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 g in 100 g	
BENZO CAINE (UNII: U3RS Y48 JW5) (BENZO CAINE - UNII:U3RS Y48 JW5)	BENZOCAINE	10 g in 100 g	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.33 g in 100 g	

Inactive Ingredients			
Ingredient Name	Strength		
ISOBUTANE (UNII: BXR49TP611)			
BUTANE (UNII: 6LV4FOR43R)			
PROPANE (UNII: T75W9911L6)			
DIPROPYLENE GLYCOL (UNII: E107L85C40)			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:0498-0021-40	85 g in 1 CAN; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/12/2018	

Part 5 of 9

TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

Product Information	
Item Code (Source)	NDC:0498-0750
Route of Administration	TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g	
BACITRACIN ZINC (UNII: 89 Y4M234ES) (BACITRACIN - UNII:58 H6 RWO 52I)	BACITRACIN	400 [iU] in 1 g	
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMYCIN	3.5 mg in 1 g	

Inactive Ingredients	
Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

l	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:0498-0750-36	0.5 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part333B	09/19/2018		

Part 6 of 9

ALCOHOL WIPE

isopropyl alcohol swab

Product Information

Item Code (Source) NDC:0498-0143

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

Ingredient Name Strength

WATER (UNII: 059QF0KO0R)

Packaging

l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1	NDC:0498-0143-04	0.4 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/18/2018	

Part 7 of 9

MIRALAC

calcium carbonate tablet

Product Information

Item Code (Source)	NDC:0498-0303
Route of Administration	ORAL.

Active Ingredient/Active Moiety

Ingredient Name	Basis of	Strength
ingredient Name	Strength	Strength

420 mg

Inactive Ingredients			
Ingredient Name	Strength		
MAGNESIUM STEARATE (UNII: 70097M6I30)			
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)			
SORBITOL (UNII: 506T60A25R)			
STARCH, CORN (UNII: O8232NY3SJ)			

Product Characteristics				
Color	white	Score	2 pieces	
Shape	ROUND	Size	11mm	
Flavor	MINT	Imprint Code	FR8	
Contains				

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part331	02/22/2012		

Part 8 of 9

ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

Product Information	
Item Code (Source)	NDC:0498-0501
Route of Administration	TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL		

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

l	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:0498-0501-00	1.4 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	12/22/20 17		

Part 9 of 9

AYPANAL EX

acetaminophen tablet

Product Information	
Item Code (Source)	NDC:0498-2110
Route of Administration	ORAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ACETAMINO PHEN (UNII: 36209 ITL9D) (ACETAMINO PHEN - UNII: 36209 ITL9D)	ACETAMINOPHEN	500 mg		

Inactive Ingredients				
Ingredient Name	Strength			
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)				
STARCH, CORN (UNII: O8232NY3SJ)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
PO VIDO NE (UNII: FZ989 GH94E)				
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)				

Product Characteristics				
Color	white	Score	no score	
Shape	ROUND	Size	12mm	
Flavor		Imprint Code	FR1	
Contains				

Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC:0498-2110-01 2	in 1 PACKET; Type 0: Not a Combination Product		
Marketing Info	rmation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	01/02/2017	
ore monograph nor mar	parto-45	01/02/201/	
OTC monograph not finat	parto45	01/02/201/	
OTC monograph noctimat	parts45	0 102/2017	
Marketing Info	'	0 102/2017	
U 1	'	Marketing Start Date	Marketing End Date

Labeler - Honeywell Safety Products USA, INC (079287321)

Establishment			
Name	Address	ID/FEI	Business Operations
James Alexander		040756421	manufacture(0498-3334)

Establishment			
Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, INC		079287321	pack(0498-4339)

Establishment			
Name	Address	ID/FEI	Business Operations
Ultra Seal Corporation		085752004	manufacture(0498-2110, 0498-0303)

Establishment			
Name	Address	ID/FEI	Business Operations
Dixon Investments		115315822	manufacture(0498-0031, 0498-0021)

Establishment			
Name	Address	ID/FEI	Business Operations
Water-Jel Technologies		155522589	manufacture(0498-0750)

Establishment			
Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, Inc.		167518617	manufacture(0498-0100)

Establishment				
Name	Address	ID/FEI	Business Operations	
Changzhou Maokang Medical		421317073	manufacture(0498-0143, 0498-0501)	